CLAIMS

- A process for the preparation of a dry powder inhaler which comprises exposing, during manufacture, a dry powder inhaler optionally filed with a powder formulation, or one or
 more components thereof, to a gas at low pressure.
 - 2. A process according to claim 1 where the gas is air.
 - 3. A process according to claim 1 or 2 where the pressure is less than 200 mbar.

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- 4. A process according to any one of claims 1 to 3 where the pressure is less than 100 mbar.
- 5. A process according to any one of claims 1 to 4 where the inhaler is Turbuhaler.

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- 6. A process according to any one of claims 1 to 5 where the powder formulation comprises one or more drugs and one or more pharmaceutically acceptable carriers or excipients.
- 7. A process according to any one of claims 1 to 5 where the drug is mometasone (e.g. as furoate), ciclesonide, zoticasone, flumoxonide, fluticasone (e.g. as 17-propionate) budesonide, salmeterol (e.g. as xinafoate) and formoterol (e.g. as furnarate dihydrate) or ttiotropium bromide.
- 25 8. A process according to any one of claims 1 to 5 where the drugs are combinations of fluticasone propionate/salmeterol xinafoate, ciclesonide/formoterol fumarate dihydrate, mometasone furoate/formoterol fumarate dihydrate, budesonide/formoterol fumarate dihydrate and tiotropium bromide/formoterol fumarate dihydrate.

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9. A process according to any one of claims 1 to 5 where the drugs are combinations of, budesonide and formoterol furnarate dihydrate.

- 10. A process for reducing electrostatic charges from one or more inhaler components, or a complete inhaler, optionally filed with a powder formulation, the process comprising:
 - i) placing one or more inhaler components, or a complete inhaler, optionally filed with a powder formulation in a chamber,
 - ii) reducing the pressure of gas in the chamber,
 - iii) returning the pressure to atmospheric pressure
 - iv) optionally repeating steps ii) and iii)

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10 11. A dry powder inhaler prepared according to the process of any one of claims 1 to 10.